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112110	-	1100		minutes
NEWS	3	AUG	18	COMPENDEX indexing changed for the Corporate Source
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				CAS REGISTRY
NEWS	7	SEP	11	WPIDS, WPINDEX, and WPIX now include Japanese FTERM
				thesaurus
NEWS	8	OCT	21	Derwent World Patents Index Coverage of Indian and
				Taiwanese Content Expanded
NEWS	9	OCT	21	Derwent World Patents Index enhanced with human
				translated claims for Chinese Applications and
				Utility Models
NEWS		NOV		Addition of SCAN format to selected STN databases
NEWS		NOV		Annual Reload of IFI Databases
NEWS				FRFULL Content and Search Enhancements
NEWS	13	DEC	01	DGENE, USGENE, and PCTGEN: new percent identity
				feature for sorting BLAST answer sets
NEWS	14	DEC	02	Derwent World Patent Index: Japanese FI-TERM
				thesaurus added
NEWS	15	DEC	02	PCTGEN enhanced with patent family and legal status
				display data from INPADOCDB
NEWS	16	DEC	02	USGENE: Enhanced coverage of bibliographic and
				sequence information
NEWS	17	DEC	21	New Indicator Identifies Multiple Basic Patent
				Records Containing Equivalent Chemical Indexing
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NEWS	10	JAN	0.5	Needs, Quickly and Conveniently
				Annual Reload of MEDLINE database STN Express Maintenance Release, Version 8.4.2, Is
NEWS	20	FEB	Тρ	Now Available for Download
NEWS	21	FEB	16	Derwent World Patents Index (DWPI) Revises Indexing
MEMO	41	rab	10	of Author Abstracts
NEWS	22	FEB	16	New FASTA Display Formats Added to USGENE and PCTGEN
NEWS		FEB		INPADOCDB and INPAFAMDB Enriched with New Content
MEMO	23	200	10	THEADOCDD GIRG INFAFABIDD BILLICHEG WICH NEW CONCENT

<12/04/2007> Erich Leese

and Features

NEWS 24 FEB 16 INSPEC Adding Its Own IPC codes and Author's E-mail

NEWS EXPRESS FEBRUARY 15 10 CURRENT WINDOWS VERSION IS V8.4.2, AND CURRENT DISCOVER FILE IS DATED 15 JANUARY 2010.

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=> file reg COST IN U.S. DOLLARS

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FULL ESTIMATED COST 0.22 0.22
FILE 'REGISTRY' ENTERED AT 14:56:13 ON 08 MAR 2010

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ENTRY

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STRUCTURE FILE UPDATES: 7 MAR 2010 HIGHEST RN 1208070-84-1 DICTIONARY FILE UPDATES: 7 MAR 2010 HIGHEST RN 1208070-84-1

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```
11 12 13 14 15 17 19 21 24 26 32 34 35 45 46 47 48 49
ring nodes :
1 2 3 4 5 6 7 8 9 10 27 28 29 30 31 37 38 39 40 41 42
chain bonds :
1-26 2-24 3-21 4-19 7-17 8-12 9-11 12-14 12-13 14-15 14-28 27-32 29-34
30-35 31-39 37-49 38-48 40-45 41-46 42-47
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 5-7 6-10 7-8 8-9 9-10 27-28 27-31 28-29 29-30
30-31 37-38 37-42 38-39 39-40 40-41 41-42
exact/norm bonds :
1-26 2-24 3-21 4-19 5-7 6-10 7-8 7-17 8-9 9-10 9-11 12-14 12-13 14-28
27-31 27-32 29-30 29-34 30-31 30-35 31-39 37-49 38-48 40-45 41-46 42-47
exact bonds :
8-12 14-15 27-28 28-29
normalized bonds :
1-2 1-6 2-3 3-4 4-5 5-6 37-38 37-42 38-39 39-40 40-41 41-42
isolated ring systems :
containing 27 : 37 :
```

G1:C,H

G2:H, Ak

G3:H, X, Ak, NO2, C

chain nodes :

G4:H,OH,Ak,MeO,EtO,n-PrO,i-PrO,n-BuO,i-BuO,s-BuO,t-BuO,C,O

<12/04/2007>

Erich Leese

G5:H, Ak, O, NO2

G6:CH3,Et

G7:H, CH3

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 17:CLASS 19:CLASS 21:CLASS

24:CLASS 26:CLASS 27:Atom 28:Atom 29:Atom 30:Atom 31:Atom 32:CLASS 34:CLASS 35:CLASS 37:CLASS 38:CLASS 39:CLASS 40:CLASS 41:CLASS 42:CLASS 45:CLASS

46:CLASS 47:CLASS 48:CLASS 49:CLASS

L1 STRUCTURE UPLOADED

=> d 11

- G1 C,H
- G2 H, Ak
- G3 H, X, Ak, NO2, C
- G4 H, OH, Ak, MeO, EtO, n-PrO, i-PrO, n-BuO, i-BuO, s-BuO, t-BuO, C,O
- G5 H, Ak, O, NO2
- G6 Me,Et
- G7 H, Me

Structure attributes must be viewed using STN Express query preparation.

=> s 11 sss

SAMPLE SEARCH INITIATED 14:57:28 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED -1 TO ITERATE

100.0% PROCESSED 1 ITERATIONS 1 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
PROJECTED ITERATIONS: 1 TO 80
PROJECTED ANSWERS: 1 TO 80

L2 1 SEA SSS SAM L1

=> d scan

- L2 1 ANSWERS REGISTRY COPYRIGHT 2010 ACS on STN
- IN 2H-1-Benzopyran-3-carboxamide, N-(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4-yl)-4-methyl-2-oxo-
- MF C22 H19 N3 O4

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

=> s 11 full

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FULL SCREEN SEARCH COMPLETED - 21 TO ITERATE

100.0% PROCESSED 21 ITERATIONS 14 ANSWERS

SEARCH TIME: 00.00.01

L3 14 SEA SSS FUL L1

=> file caplus

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FILE COVERS 1907 - 8 Mar 2010 VOL 152 ISS 11
FILE LAST UPDATED: 7 Mar 2010 (20100307/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Dec 2009
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Dec 2009

CAplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2009.

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 13 full L4 4 L3

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L4 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 2009:769551 CAPLUS

DOCUMENT NUMBER: 151:70320

TITLE: Method using lifespan-altering compounds for altering

the lifespan of eukaryotic organisms, and screening

for such compounds

INVENTOR(S): Goldfarb, David Scott
PATENT ASSIGNEE(S): University of Rochester, USA

SOURCE: U.S. Pat. Appl. Publ., 57pp.

CODEN: USXXCO
DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 20 PATENT INFORMATION:

PAT	TENT					KIND DATE					LICAT:								
US	2009						0625			2008-3		20081222							
US	2009	0163	545		A1		2009	0625		US :	2008-3	3416	20081222						
US	2009	0163	545		A1		2009	0625		US :	2008-3	3416	20081222						
US	2009	0163	545		A1		2009	0625		US :	2008-3	3416	20081222						
US	2009	0163	545		A1	2009	0625		US :	2008-3	3416	20081222							
US	2009	0163	545		A1	2009	0625		US :	2008-3	3416	20081222							
US	US 20090163545						2009	0625		US :	2008-3	3416	20081222						
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US	2009	0163	545		A1		2009	0625		US :	2008-3	3416	20081222						
US	2009	0163	545		A1		2009	0625		US :	2008-3	3416	15		20081222				
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US	2009	0163	545		A1		20090625 US 2008-341615								20081222				
	2009				A1		2009	0625		US :	2008-3	3416							
US	2009	0163	545		A1										2	0081	222		
	US 20090163545						2009	0625		US :	2008-3	3416	20081222						
	US 20090163545						2009	0625		US :	2008-3	3416	20081222						
	US 20090163545							0625			2008-3		20081222						
US	2009	0163	545		A1		2009	0625			2008-3		20081222 20081222						
US	2009	0163	545		A1		2009	0625											
	2009				A1			0625						20081222					
	2009							0625					20081222						
WO	2009	0863	03				2009	0709		WO :	2008-ī	JS88	20081222						
WO	2009							1230											
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ORITY	Y APP	LN.	INFO	. :						US :	2007-	1636	2P	1	P 2	0071	2.21		

PRIORITY APPLN. INFO.:

US 2007-16362P P 20071221 US 2008-23801P P 20080125 US 2008-341615 20081222

- AB The invention discloses a method for altering the lifespan of a eukaryotic organism. The method comprises the steps of providing a lifespan-altering compound, and administering an effective amount of the compound to a eukaryotic organism, such that the lifespan of the organism is altered. In one embodiment, the compound is identified using the DeaD assay. [This abstract record is one of 20 records for this document necessitated by the large number of index entries required to fully index the document and publication system constraints.]
- IT 301681-84-5 RL: PAC (Pharmacological activity); BIOL (Biological study) (method using lifespan-altering compds for altering lifespan of eukaryotic organisms, and screening for such compds.)
- RN 301681-84-5 CAPLUS
 CN 2H-1-Bencopyran-3-carboxamide, N-(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl1H-pyrazol-4-yl)-2-oxo- (CA INDEX NAME)

OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

L4 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 2004:490719 CAPLUS

DOCUMENT NUMBER: 141:38530

TITLE: Preparation of coumarincarboxamides as TGF-β

inhibitors

INVENTOR(S): Xu, Shiping; Chen, Xiaoguang; Xu, Song; Li, Lanmin; Xie, Longfei; Li, Hongvan; Li, Yan; Cheng, Guifang

PATENT ASSIGNEE(S): Institute of Materia Medica, Chinese Academy of

Medical Sciences, Peop. Rep. China

SOURCE: PCT Int. Appl., 79 pp.

CODEN: PIXXD2
DOCUMENT TYPE: Patent

LANGUAGE: Chinese
FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

					KIND DATE															
									WO 2003-CN1046											
	W:	ΑE,	AG,	AL,	AM,	AT,	ΑU,	AZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,			
		CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,			
		GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KΡ,	KR,	ΚZ,	LC,	LK,	LR,			
		LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NI,	NO,	NZ,	OM,			
		PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	TJ,	TM,	TN,			
		TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW						
	RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	AZ,			
		BY,	KG,	ΚZ,	MD,	RU,	ΤJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,			
		ES,	FI,	FR,	GB,	GR,	HU,	IE,	IT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,			
		TR,					CI,											TG		
	CN 1506359								CN 2002-155525											
	CA 2508573								CA 2003-2508573											
							AU 2003-289641													
EP					A1 20050907			EP 2003-776786												
	R:						ES,										PT,			
							RO,													
BR	BR 2003016595						2005	1004	BR 2003-16595						20031205					
JP	JP 2006512328					T 20060413					JP 2004-555972 CN 2003-80109516						20031205			
CN	CN 1829506						2006	0906		CN 2	003-	8010	9516		2	0031	205			
CN	1004	8850	4		C		2009	0520												
	RU 2361870						2009													
										IN 2005-DN2384										
						A1 20060706														
PRIORIT:	RIORITY APPLN. INFO									CN 2002-155525										
										WO 2	003-	CN10	46		W 2	0031	205			

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT

OTHER SOURCE(S): MARPAT 141:38530

GI

AB Title compds. I [R3 = H, carboxy, ethoxycarbonyl, CONH(CH2)3CO2H, 5-phenyloxadiazol-Z-yl, CONHRC, etc.; R2 = carboxylic acid, benzoylamino, nicotinoylamino, Ph, substituted Ph, etc.; R4 = H, CONHRI; R1 = carboxylic acid, benzoylamino, isonicotinoylamino, Ph, substituted Ph, etc; R5 = H, alkyl; R6 = H, alkyl, halo, NO2 etc.; R7 = H, OH, alkyl, alkoxy, carboxyalkoxy, etc.; R8 = H, alkyl, alkoxy, NO2], useful as TGF-B inhibitors, are prepared Thus, 3-carboxy-6-ethyl-7-methoxycoumarin was chlorinated with SCC12 followed by reaction with 4-aminosalicylic acid in pyridine gave 3-(3-hydroxy-4-carboxyphenylaminocarbonyl)-6-ethyl-7-methoxycoumarin. The invention also discloses the drug composition comprising said compariand derivs., and the use for especially kidney protection, treatment for hypertension, cerebrovascular and cardiovascular diseases, diabetes II,

- tumor, precancerous lesion and dropsy.

 17 704881-98-1P 704882-10-0P 704882-14-4P

 RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (preparation of coumarincarboxamides as TGF-β inhibitors)
- RN 704881-98-1 CAPLUS

Ι

CN 2H-1-Benzopyran-3-carboxamide, N-(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4-yl)-7-methoxy-2-oxo- (CA INDEX NAME)

- RN 704882-10-0 CAPLUS
- CN 2H-1-Benzopyran-3-carboxamide, 6-bromo-N-(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4-yl)-7-methoxy-2-oxo- (CA INDEX NAME)

RN 704882-14-4 CAPLUS

CN 2H-1-Benzopyran-3-carboxamide, N-(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4-yl)-6-hexyl-7-methoxy-2-oxo- (CA INDEX NAME)

OS.CITING REF COUNT: 3 THERE ARE 3 CAPLUS RECORDS THAT CITE THIS RECORD (3 CITINGS)

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 1988:630878 CAPLUS DOCUMENT NUMBER: 109:230878

ORIGINAL REFERENCE NO.: 109:38185a,38188a

Application of the Knoevenagel condensation to TITLE:

4-acetamidophenazone derivatives

AUTHOR(S): El-Kerdawy, M. M.; Farghaly, A. M.; Massoud, M. A.

CORPORATE SOURCE: Fac. Pharm., Mansoura Univ., Mansoura, Egypt SOURCE: Indian Journal of Chemistry, Section B: Organic

Chemistry Including Medicinal Chemistry (1987),

ΙI

26B(12), 1189-91

CODEN: IJSBDB; ISSN: 0376-4699

DOCUMENT TYPE: Journal LANGUAGE:

English

OTHER SOURCE(S): CASREACT 109:230878

GI

AB Antipyrines I (R1 = CO2Et, cyano) were treated with carbonyl compds. (e.g., PhCH:CHCHO, retinal, Me2CO, MeCOEt, cyclohexanone) and piperidine to give the resp. condensation products II. Coumarin derivative III was obtained from I (R1 = CO2Et) and 2,4-(HO)2C6H3CHO.

117665-36-8P 117665-37-9P 117665-42-6P 117665-43-7P

RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

RN 117665-36-8 CAPLUS

CN 2H-1-Benzopyran-3-carboxamide, N-(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4-vl)-7-hydroxy-2-oxo- (CA INDEX NAME)

- RN 117665-37-9 CAPLUS
- CN 2H-1-Benzopyran-3-carboxamide, N-(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4-yl)-6-nitro-2-oxo- (CA INDEX NAME)

- RN 117665-42-6 CAPLUS
- CN 2H-1-Benzopyran-3-carboxamide, 7-(acetyloxy)-N-(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4-yl)-2-oxo- (CA INDEX NAME)

- RN 117665-43-7 CAPLUS
- CN 2H-1-Benzopyran-3-carboxamide, N-(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4-yl)-4-methyl-2-oxo- (CA INDEX NAME)

- OS.CITING REF COUNT: 2
- THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD
 (2 CITINGS)

L4 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2010 ACS on STN ACCESSION NUMBER: 1968:402804 CAPLUS

DOCUMENT NUMBER: 69:2804

ORIGINAL REFERENCE NO.: 69:535a,538a

Coumarin derivatives of pharmaceutical interest TITLE: AUTHOR(S):

Selleri, R.; Orzalesi, G.; Caldini, O.; Spano, R.;

Ferretti, G.

CORPORATE SOURCE: Soc. Italo-Britannica L. Manetti, H. Roberts Co.,

Florence, Italy

SOURCE: Bollettino Chimico Farmaceutico (1967), 106(10), 680-7

CODEN: BCFAAI; ISSN: 0006-6648

DOCUMENT TYPE: Journal LANGUAGE: Italian

GT For diagram(s), see printed CA Issue.

Compds. of the general formula I are prepared A mixture of 0.05M ΔB

6-carbomethoxycoumarin-3-carboxylic acid and 200 ml. SOC12 is refluxed 2-4 hrs. and the acid chloride treated with 0.1 mole NH3 to give Me 3-carbamoylcoumarin-6-carboxylate, m. 274° (MeOH). Similarly prepared are the following I (R, R1, R2, and m.p. given): H, H, Et,

259° (EtOH); H, Pr, Me, 183° (MeOH); H, Pr, Et, 166° (EtOH); H, Bu, Me, 181° (MeOH); H, Bu, Et, 142° (EtOH); Me, Me, Me, 178° (MeOH); Me, Me, Et, 152° (EtOH); Et, Et, Me,

154° (MeOH); Et, Et, Et, 133° (EtOH); (NRR1 =) pyrrolidino, -, Me, 155° (MeOH); (NRR1 =) pyrrolidino, -, Et, 184° (EtOH); (NRR1 =) morpholino, -, Me, 238° (MeOH); (NRR1 =)

morpholino, -, Et, 185° (EtOH); H, Ph, Me, 208° (MeOH); H, Ph, Et, 206° (HOAc); H, PhCH2, Me, 202° (MeOH); H, PhCH2, Et, 187° (EtOH); H, p-EtOC6H4, Me, 227° (MeOH); H,

p-EtOCC6H4, Et, 213° (EtOH); H, p-EtO2CC6H4, Me, 246° (HOAc); H, p-EtO2CC6H4, Et, 235° (HOAc); H, p-MeSO2C6H4, Me, 297° (MeOH); H, p-MeSO2C6H4, Et, 266° (EtOH); H, CONH2, Me, 256° (MeOH); and H, CONH2, Et, 250° (EtOH). Also prepared were I (R = H,

R1 = 1-phenyl-2, 3-dimethyl-5-oxo-3-pyrazolin-4-yl, R2 = Me) (II), m. 230° (HOAc); and I (R = H, R1 = 1-phenyl-2,3-dimethyl-5-oxo-3-pyrazolin-4-yl, R1 = Et) (III), m.

191° (HOAc). II and III demonstrate analgesic activity in mice. (R = R1 = Et, R2 = Me) increases respiration and blood pressure in

16409-08-8P 16409-09-9P

> RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)

RN 16409-08-8 CAPLUS

rabbits.

CN 2H-1-Benzopyran-6-carboxylic acid,

3-[[(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4v1)amino|carbonv1|-2-oxo-, methvl ester (CA INDEX NAME)

RN 16409-09-9 CAPLUS

CN 2H-1-Benzopyran-6-carboxylic acid,

3-[[(2,3-dihydro-1,5-dimethyl-3-oxo-2-phenyl-1H-pyrazol-4-yl)amino]carbonyl]-2-oxo-, ethyl ester (CA INDEX NAME)

OS.CITING REF COUNT: 2 THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD (3 CITINGS)

=> d his

(FILE 'HOME' ENTERED AT 14:55:56 ON 08 MAR 2010)

FILE 'REGISTRY' ENTERED AT 14:56:13 ON 08 MAR 2010

L1 STRUCTURE UPLOADED L2 1 S L1 SSS

L3 14 S L1 FULL

FILE 'CAPLUS' ENTERED AT 14:57:51 ON 08 MAR 2010 L4 $$\rm 4\ S\ L3\ FULL$$

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COST ÎN U.S. DOLLARS

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